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Selenium and outcome in heart failure

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Aims

Severe deficiency of the essential trace element selenium can cause myocardial dysfunction although the mechanism at cellular level is uncertain. Whether, in clinical practice, moderate selenium deficiency is associated with worse symptoms and outcome in patients with heart failure is unknown.

Methods and results

BIOSTAT-CHF is a multinational, prospective, observational cohort study that enrolled patients with worsening heart failure. Serum concentrations of selenium were measured by inductively coupled plasma mass spectrometry. Primary endpoint was a composite of all-cause mortality and hospitalization for heart failure; secondary endpoint was all-cause mortality. To investigate potential mechanisms by which selenium deficiency might affect prognosis, human cardiomyocytes were cultured in absence of selenium, and mitochondrial function and oxidative stress were assessed. Serum selenium concentration (deficiency) was $<70 \mu\text{g/L}$ in 485 (20.4%) patients, who were older, more often women, had worse New York Heart Association class, more severe signs and symptoms of heart failure and poorer exercise capacity (6-min walking test) and quality of life (Kansas City Cardiomyopathy Questionnaire). Selenium deficiency was associated with higher rates of the primary endpoint [hazard ratio (HR) 1.23; 95% confidence interval (CI) 1.06–1.42] and all-cause mortality (HR 1.52; 95% CI 1.26–1.86). In cultured human cardiomyocytes, selenium deprivation impaired mitochondrial function and oxidative phosphorylation, and increased intracellular reactive oxygen species levels.

Conclusions

Selenium deficiency in heart failure patients is independently associated with impaired exercise tolerance and a 50% higher mortality rate, and impaired mitochondrial function *in vitro*, in human cardiomyocytes. Clinical trials are needed to investigate the effect of selenium supplements in patients with heart failure, especially if they have low plasma concentrations of selenium.

Keywords

Selenium • Malnutrition • Heart failure • Mitochondrial function • Cardiomyocytes • All-cause mortality

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[†]<https://cordis.europa.eu/project/rcn/94179/reporting/en>

Introduction

An aberrant equilibrium of circulating molecules like minerals and trace elements (for example iron, iodine and zinc) in the patients' blood is closely involved in the development and progression of heart failure.^{1–4} It was shown that up to 50% of patients with heart failure suffer from some form of malnutrition, like micronutrient insufficiencies.^{5,6}

Selenium is an essential micronutrient and sufficient concentrations are essential for numerous biological functions including thyroid hormone metabolism, antioxidant defenses, the immune system and in certain types of cancer.^{7,8} Severe selenium deficiency in humans is associated with a rare but fatal form of dilated cardiomyopathy that is restricted to specific geographic regions (Keshan disease)⁹ that have a very low amount of selenium in the soil and therefore in food. Keshan disease is reversible with selenium supplementation.¹ Selenium soil content is very variable and this affects dietary selenium intake. For example, intakes are high in Venezuela, Canada, the USA, and Japan (>100 µg/day), and much lower in some parts of Europe (~40 µg/day).⁷ Observational studies in the general population^{7,10} suggest that selenium deficiency might be common, however data on serum selenium concentrations in heart failure are scarce.^{11–13}

On a molecular level selenium is incorporated into, and is essential for the enzymatic activity of 25 different selenoproteins.¹³ Selenium depletion deprives the cell of its ability to synthesize enzymatically active selenoproteins, but the effects on mitochondrial function, biogenesis and oxidative stress in human cardiomyocytes are unknown.

Accordingly, we conducted both *in vitro* experiments on the effects of selenium deprivation on human cardiomyocytes and investigated associations between serum concentrations of selenium and the clinical characteristics and outcomes of a large cohort of patients with heart failure.

Materials and methods

Abbreviated methods are displayed below, for the full materials and methods section, see online supplementary *Methods S1*.

BIOSTAT-CHF patient population

We used information and samples from the BIOSAT-CHF index cohort, which consists of 2516 patients with heart failure from 69 centres in 11 European countries.^{14–17} Inclusion criteria included: age >18 years, symptoms of new-onset or worsening heart failure and at least one of the following: a left ventricular ejection fraction of ≤40%, a plasma B-type natriuretic peptide of >400 pg/mL or a plasma N-terminal pro-B-type natriuretic peptide (NT-proBNP) of >2000 pg/mL. Patients could only be included if their treatment did not conform to guideline recommendations [i.e. ≤50% of angiotensin-converting enzyme inhibitors (ACEi)/angiotensin receptor blockers (ARB) and/or beta-blockers] and were anticipated to have treatment with ACEi/ARBs and/or beta-blockers initiated or up-titrated. The BIOSAT-CHF study was approved by the medical ethics committee for each centre and complies with the Declaration of Helsinki. All participants provided written informed consent prior to any study-related procedures.

For additional details on the BIOSAT-CHF study definitions, measurements and outcome analyses and statistical analysis, see online supplementary *Methods S1*.

Serum selenium determination

Selenium was measured using a validated inductively coupled plasma mass spectrometry (ICP-MS) method with a lower limit of quantitation of 20 µg/L serum. The ⁷⁸Se isotope concentration was measured using a Varian 820-MS ICP mass spectrometer. Reference ranges for serum selenium vary between countries because of differences in dietary sources of selenium. For caucasians of European ancestry, adult reference ranges are between 70 and 130 µg/L.¹⁸ Deficiency of selenium was therefore set at serum levels <70 µg/L.

For additional details on serum selenium determination, see online supplementary *Methods S1*.

Cell culture, cardiomyocyte differentiation, selenium depletion and cardiac stress model

Differentiation of HUES9 human pluripotent stem cells (hPSC, Harvard Stem Cell Institute) to cardiomyocytes was achieved as described previously.^{19–22} For the experiments, cells were grown either in CDM3 medium supplemented with 100 nM sodium selenite²³ (Control; S5261, Sigma-Aldrich) or in CDM3 medium without added selenium (selenium deficient cells) for 2 weeks.

For additional details on the functional characterisation of selenium depletion of cardiomyocytes (protein extraction and western blot, RNA extraction and qRT-PCR, immunofluorescence, reactive oxygen species detection and Seahorse experiments), see online supplementary *Methods S1* and *Table S1*.

Statistical analyses

Differences between clinical characteristics across quartiles of selenium levels were compared using one-way analysis of covariance (ANOVA), the Kruskal–Wallis test or the Chi square test where appropriate. Multivariable logistic regression was used to investigate independent associations with selenium deficiency. The association with outcome was investigated using Kaplan–Meier curves and the log-rank test. For multivariable analyses, Cox regression analysis was performed on selenium deficiency and standardized selenium levels, correcting for relevant clinical confounders independently associated with selenium levels (online supplementary *Table S2*) and the BIOSAT-CHF risk models, which were previously published.¹⁵ The baseline model included, age, daily protein intake, country, haemoglobin, C-reactive protein, high-density lipoprotein cholesterol, albumin, NT-proBNP, sodium and estimated glomerular filtration rate (eGFR by the Chronic Kidney Disease Epidemiology Collaboration equation), all independently associated with selenium levels. The BIOSAT-CHF risk model for predicting mortality included, age, blood urea nitrogen, NT-proBNP, haemoglobin and the use of a beta-blocker at time of inclusion. The BIOSAT-CHF risk model for predicting mortality or heart failure hospitalization included age, NT-proBNP, haemoglobin, the use of a beta-blocker at time of inclusion, a heart failure hospitalization in the year before inclusion, peripheral oedema, systolic blood pressure, high-density lipoprotein cholesterol and sodium.

Experimental groups consisted of at least three independent biological replicates and technical duplicates were used. Data shown are

Table 1 Baseline characteristics

	Normal selenium ($\geq 70 \mu\text{g/L}$)	Selenium-deficient ($< 70 \mu\text{g/L}$)	P-value
<i>n</i>	1897	485	
Demographics			
Age (years)	67.8 (12.0)	72.6 (11.3)	<0.001
Female sex	456 (24.0%)	158 (32.6%)	<0.001
BMI (kg/m^2)	28.0 (5.5)	27.5 (5.5)	0.10
Protein intake (g/day)	55.8 (11.5)	52.2 (9.9)	<0.001
Ischaemic aetiology (%)	870 (46.7%)	215 (44.9%)	0.47
LVEF (%)	30.5 (9.9)	32.9 (12.7)	<0.001
HF subgroup			<0.001
HFrEF	1430 (83.6%)	303 (71.8%)	
HFmrEF	194 (11.3%)	73 (17.3%)	
HFpEF	86 (5.0%)	46 (10.9%)	
NYHA class			<0.001
I	198 (11.9%)	27 (6.2%)	
II	896 (54.0%)	201 (46.4%)	
III	512 (30.9%)	179 (41.3%)	
IV	52 (3.1%)	26 (6.0%)	
Systolic BP (mmHg)	124.5 (21.5)	124.6 (22.7)	0.93
Diastolic BP (mmHg)	75.3 (13.2)	73.4 (12.9)	0.005
Heart rate (bpm)	79.6 (19.5)	80.1 (18.9)	0.63
Smoking			0.074
No	670 (35.4%)	196 (40.5%)	
Past	943 (49.8%)	229 (47.3%)	
Current	282 (14.9%)	59 (12.2%)	
Signs and symptoms			
Extent of peripheral oedema			<0.001
Not present	706 (44.9%)	100 (24.9%)	
Ankle	456 (29.0%)	121 (30.2%)	
Below knee	320 (20.4%)	121 (30.2%)	
Above knee	90 (5.7%)	59 (14.7%)	
Elevated JVP	384 (30.0%)	138 (46.9%)	<0.001
Hepatomegaly	252 (13.3%)	81 (16.8%)	0.049
Orthopnoea	605 (32.0%)	215 (44.4%)	<0.001
Pulmonary congestion	943 (51.3%)	291 (61.0%)	<0.001
Co-morbidities			
Anaemia	567 (33.2%)	222 (47.9%)	<0.001
Atrial fibrillation	839 (44.2%)	238 (49.1%)	0.056
Diabetes mellitus	599 (31.6%)	170 (35.1%)	0.14
COPD	313 (16.5%)	98 (20.2%)	0.054
CKD	487 (25.7%)	170 (35.1%)	<0.001
Hypertension	1178 (62.1%)	310 (63.9%)	0.46
Peripheral arterial disease	197 (10.4%)	56 (11.5%)	0.46
Stroke	167 (8.8%)	50 (10.3%)	0.30
PCI	429 (22.6%)	98 (20.2%)	0.25
CABG	320 (16.9%)	90 (18.6%)	0.38
Medication			
Loop diuretics	1889 (99.6%)	482 (99.4%)	0.57
ACEi/ARB	1391 (73.3%)	333 (68.7%)	0.040
Aldosterone antagonist	1031 (54.3%)	243 (50.1%)	0.094
Beta-blocker	1608 (84.8%)	382 (78.8%)	0.001
Antiplatelets	983 (51.8%)	252 (52.0%)	0.96

Table 0 (Continued)

	Normal selenium ($\geq 70 \mu\text{g/L}$)	Selenium-deficient ($< 70 \mu\text{g/L}$)	P-value
Laboratory			
Haemoglobin (g/dL)	13.4 (1.8)	12.4 (1.9)	<0.001
Ferritin ($\mu\text{g/L}$)	107 (52–199)	76 (41–159)	<0.001
Transferrin saturation (%)	17.7 (11.7–25.6)	13.3 (9.0–21.2)	<0.001
Albumin (g/L)	33.1 (8.6)	28.9 (8.6)	<0.001
C-reactive protein (mg/L)	12.1 (5.1–24.2)	17.6 (8.4–32.3)	<0.001
Total cholesterol (mmol/L)	4.2 (3.4–5.1)	3.8 (3.1–4.6)	<0.001
HDL cholesterol (mmol/L)	1.1 (0.9–1.3)	1.0 (0.8–1.2)	0.001
LDL cholesterol (mmol/L)	2.6 (1.9–3.2)	2.2 (1.5–3.0)	<0.001
HbA1c (%)	6.3 (5.7–7.2)	6.3 (5.8–6.9)	0.96
NT-proBNP (ng/L)	3845 (2242–7681)	5506 (3045–9727)	<0.001
Sodium (mmol/L)	140 (137–142)	139 (136–141)	<0.001
Potassium (mmol/L)	4.2 (3.9–4.6)	4.3 (3.9–4.6)	0.81
Creatinine ($\mu\text{mol/L}$)	100 (82–126)	106 (83–141)	0.007
eGFR (mL/min/1.73 m ²)	61.0 (45.9–80.0)	53.7 (37.7–74.4)	<0.001

ACEi, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; BMI, body mass index; BP, blood pressure; CABG, coronary artery bypass graft; CKD, chronic kidney disease; COPD, chronic obstructive pulmonary disease; eGFR, estimated glomerular filtration rate (by Chronic Kidney Disease-Epidemiology Collaboration equation); HbA1c, glycated haemoglobin; HDL, high-density lipoprotein; HF, heart failure; HFmrEF, heart failure with mid-range ejection fraction; HFpEF, heart failure with preserved ejection fraction; HFrEF, heart failure with reduced ejection fraction; JVP, jugular venous pressure; LDL, low-density lipoprotein; LVEF, left ventricular ejection fraction; NT-proBNP, N-terminal pro-B-type natriuretic peptide; NYHA, New York Heart Association; PCI, percutaneous coronary intervention.

expressed as means \pm standard error of the mean (SEM). Differences were assessed by Student's *t*-test. A *P*-value of < 0.05 was considered statistically significant.

All tests and analyses were performed using STATA version 15.0 (StataCorp LP, College Station, TX, USA) or GraphPad Prism 7.0 (GraphPad Software, La Jolla, CA, USA).

Results

BIOSTAT-CHF baseline characteristics

Median serum selenium was $87 \mu\text{g/L}$ [interquartile range (IQR) 73–103] and 485 (20.4%) patients were selenium-deficient ($< 70 \mu\text{g/L}$). Patients with lower serum selenium concentrations were older, more often women, had worse symptoms and signs of heart failure, were more likely to have anaemia, iron deficiency and chronic kidney disease (eGFR $< 60 \text{ mL/min/1.73 m}^2$) and had lower serum albumin and cholesterol (Table 1). In multivariable analyses, country of residence, higher age, lower protein intake and lower

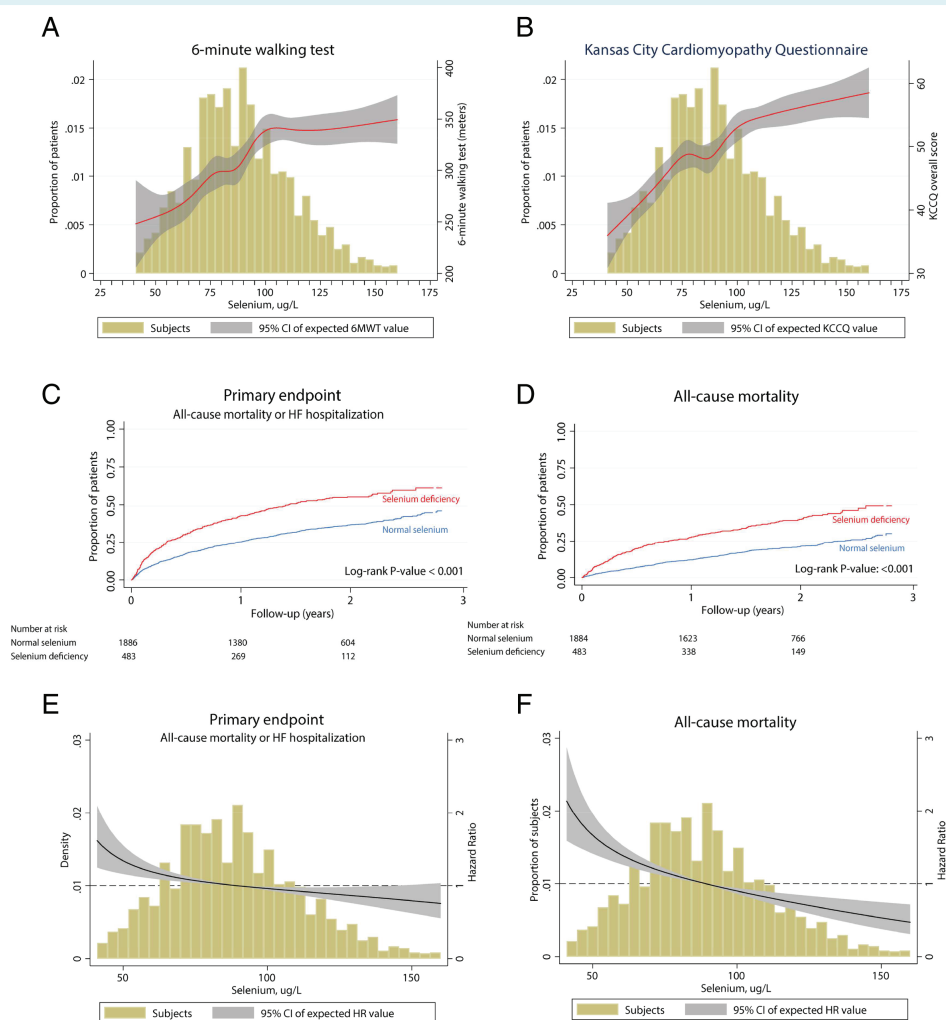


Figure 1 Selenium and clinical outcome parameters. Association of selenium levels with (A) 6-min walking test (6MWT) and (B) Kansas City Cardiomyopathy Questionnaire (KCCQ) results. Kaplan–Meier curves showing the increased risk of chronic heart failure patients with selenium deficiency (selenium $<70 \mu\text{g/L}$) for (C) the composite endpoint [all-cause mortality and hospitalization due to heart failure (HF)] and (D) all-cause mortality. Selenium levels as a continuous variable for hazard ratio (HR) plots using fractional polynomials, showing increasing HR with decreasing selenium concentrations for both the composite endpoint (E) and all-cause mortality (F). CI, confidence interval.

haemoglobin and albumin concentrations were most strongly associated with selenium deficiency (baseline model; online supplementary Table S2). There was considerable geographic variation in serum selenium concentrations (online supplementary Figure S1).

Selenium-deficient patients were less likely to complete a 6-min walking test (6MWT) (54% vs. 67%, $P < 0.001$) and those that did had a shorter 6MWT distance [151 (IQR 0–276) m vs. 225 (IQR 0–364) m; $P < 0.001$]. Selenium-deficient patients also had a lower overall score on the Kansas City Cardiomyopathy Questionnaire (KCCQ) [41 (IQR 25–57) vs. 51 (IQR 34–69)] (Figure 1A and 1B). These associations remained after multivariable correction for baseline associations with serum selenium (coefficient -30 , beta -0.07 , $P = 0.001$ and coefficient -4 , beta -0.06 , $P = 0.001$, for 6MWT and KCCQ, respectively).

Selenium deficiency associates with higher all-cause mortality and rehospitalisation for heart failure

Overall, 963 (40%) patients died or were hospitalized for heart failure during a median follow-up of 21 (IQR 16–27) months. Selenium deficiency was strongly associated [hazard ratio (HR) 1.78; 95% confidence interval (CI) 1.54–2.09] with the primary composite endpoint (Figure 1C and Table 1). This association remained significant (HR 1.21; 95% CI 1.02–1.43) after correction for baseline variables associated with serum selenium or variables included in the BIOSTAT-CHF risk model (HR 1.23; 95% CI 1.06–1.42). Similar associations were found for all-cause mortality (HR 1.52; 95% CI 1.26–1.86) (Figure 1D and Table 1). In sensitivity analyses, we analysed selenium levels as a continuous variable for HR

Table 1 Cox regression analyses

	HR (95%CI)	P-value
Combined outcome		
Univariable	1.78 (1.54–2.09)	<0.001
Baseline model ^a	1.21 (1.02–1.43)	0.028
BIOSTAT-CHF risk model ^b	1.23 (1.06–1.42)	0.007
All-cause mortality		
Univariable	2.17 (1.83–2.58)	<0.001
Baseline model ^a	1.46 (1.19–1.79)	<0.001
BIOSTAT-CHF risk model ^c	1.52 (1.26–1.86)	<0.001

CI, confidence interval; HR, hazard ratio.

^aAge, daily protein intake, country, haemoglobin, C-reactive protein, high-density lipoprotein cholesterol, albumin, N-terminal pro-B-type natriuretic peptide, sodium and estimated glomerular filtration rate (by Chronic Kidney Disease-Epidemiology Collaboration equation).

^bAge, N-terminal pro-B-type natriuretic peptide, haemoglobin, use of a beta-blocker at time of inclusion, heart failure hospitalization in the year before inclusion, peripheral oedema, systolic blood pressure, high-density lipoprotein cholesterol and sodium.

^cAge, blood urea nitrogen, N-terminal pro-B-type natriuretic peptide, haemoglobin and use of a beta-blocker at time of inclusion.

plots using fractional polynomials, which showed an increasing HR with decreasing selenium concentrations for both the composite endpoint (Figure 1E) and all-cause mortality (Figure 1F). These results were supported by the Kaplan–Meier plots of selenium concentration quartiles, showing gradual increased risk with lower concentrations (online supplementary Figure S2). Selenium concentrations were strongly inversely associated with higher rates of the primary combined outcome (HR 0.87; 95% CI 0.81–0.94) and mortality alone (HR 0.82; 95% CI 0.76–0.88) in the multivariable BIOSTAT-CHF risk model. For patients with serum selenium ≥ 70 $\mu\text{g/L}$, higher concentrations were independently associated with a better outcome (primary endpoint: HR: 0.90; 95% CI 0.82–0.99 and all-cause mortality: HR: 0.79; 95% CI 0.70–0.90).

Low selenium reduces mitochondrial functionality in human cardiomyocytes

Generated human cardiomyocytes from hPSCs (hPSC-CMs) were stained for cardiac markers, and cardiac-specific gene expression was determined for characterization purposes as described before by Hoes *et al.*²⁰

hPSC-CMs cultured for 2 weeks in selenium-free conditions showed reduced mitochondrial function (Figure 2A). Selenium depleted hPSC-CMs showed a basal levels of oxygen consumption rate of 49% ($P < 0.00001$) compared to control hPSC-CMs cultured in the presence of selenium (Figure 2A and 2B). Addition of oligomycin inhibited ATP synthase-linked respiration resulted in 48% less ATP synthase-linked respiration ($P < 0.001$) (Figure 2B). Subsequent addition of the uncoupler FCCP induced mitochondria to function at maximum capacity, which showed a 29% lower maximal capacity ($P < 0.005$) in selenium-depleted hPSC-CMs (Figure 2B). Respiration reserve capacity, an estimate of the potential bioenergetic reserve, was not significantly different between

selenium-depleted and control cells (Figure 2B). The same was true for non-mitochondrial oxygen consumption, proton leak and coupling efficiency (data not shown).

Mitochondria in selenium-depleted hPSC-CMs also had decreased expression of OXPHOS proteins [NDUFB8 (FC = 0.56), SDHB (FC = 0.55), UQCRC2 (FC = 0.61) and ATP5A (FC = 0.61) representing mitochondrial complexes 1 to 3 and 5, respectively] in hPSC-CMs with immunofluorescence (Figure 2C) and western blot (Figure 2D). To determine mitochondrial abundance, hPSC-CMs were checked for TOMM20, a mitochondrial membrane marker. Based on immunofluorescence (Figure 2E) and western blot results (Figure 2F) TOMM20 was reduced in selenium-depleted cardiomyocytes (FC = 0.72; $P = 0.004$), albeit to a lesser extent than OXPHOS. Furthermore, expression of MFN2 (mitochondrial fusion) and FIS1 (mitochondrial fission) did not change with selenium deficiency (Figure 2G).

mRNA expression levels of genes involved in (anaerobic) glycolysis or fatty acid metabolism were assessed (Figure 2G) to determine whether mitochondrial dysfunction was accompanied by a switch in metabolic substrate. Selenium-depleted hPSC-CMs had decreased expression of ACACA and ACLY, but expression of ACACB. LDHA was upregulated as the result of low selenium, but not GLUT4, PKM, HK2.

Low selenium promotes stress-induced increase of reactive oxygen species levels in human cardiomyocytes

Selenium depletion was associated with more oxidative stress. Under normal culture conditions, we found a 33% increase of reactive oxygen species (ROS) levels ($P < 0.0001$) in selenium-depleted compared to selenium supplemented cardiomyocytes. Induction of oxidative stress by adding rotenone to hPSC-CMs resulted in an additional 20% increase of ROS levels only in selenium-depleted cells ($P < 0.005$) (Figure 3A). Under oxidative stress conditions, the total increase of ROS levels between selenium-depleted and control cells was 52% ($P < 0.0001$). Addressing the expression of oxidative stress markers at a transcriptional level in selenium-depleted hPSC-CMs showed that, upon stress induction with rotenone, expression of *PGC-1 α* , *NOS2*, *AP-1* and *RELA* are increased two-fold or more ($P < 0.05$) (Figure 3B). Subsequently, under selenium replete conditions, induction of oxidative stress with rotenone, did not result in a significant induction of the stress markers *NOS2*, *AP-1* and *RELA*.

Discussion

This analysis shows that many patients with worsening heart failure have a serum selenium concentration < 70 $\mu\text{g/L}$ and that this is associated with poorer quality of life, exercise capacity and prognosis. Serum concentrations of 70–100 $\mu\text{g/L}$ appeared to have similar adverse associations, suggesting that values < 100 $\mu\text{g/L}$, found in $> 50\%$ of this cohort, might be considered abnormal.^{24,25} *In vitro* experiments on cultured human cardiac myocytes suggest that selenium deficiency disrupts mitochondrial electron transport

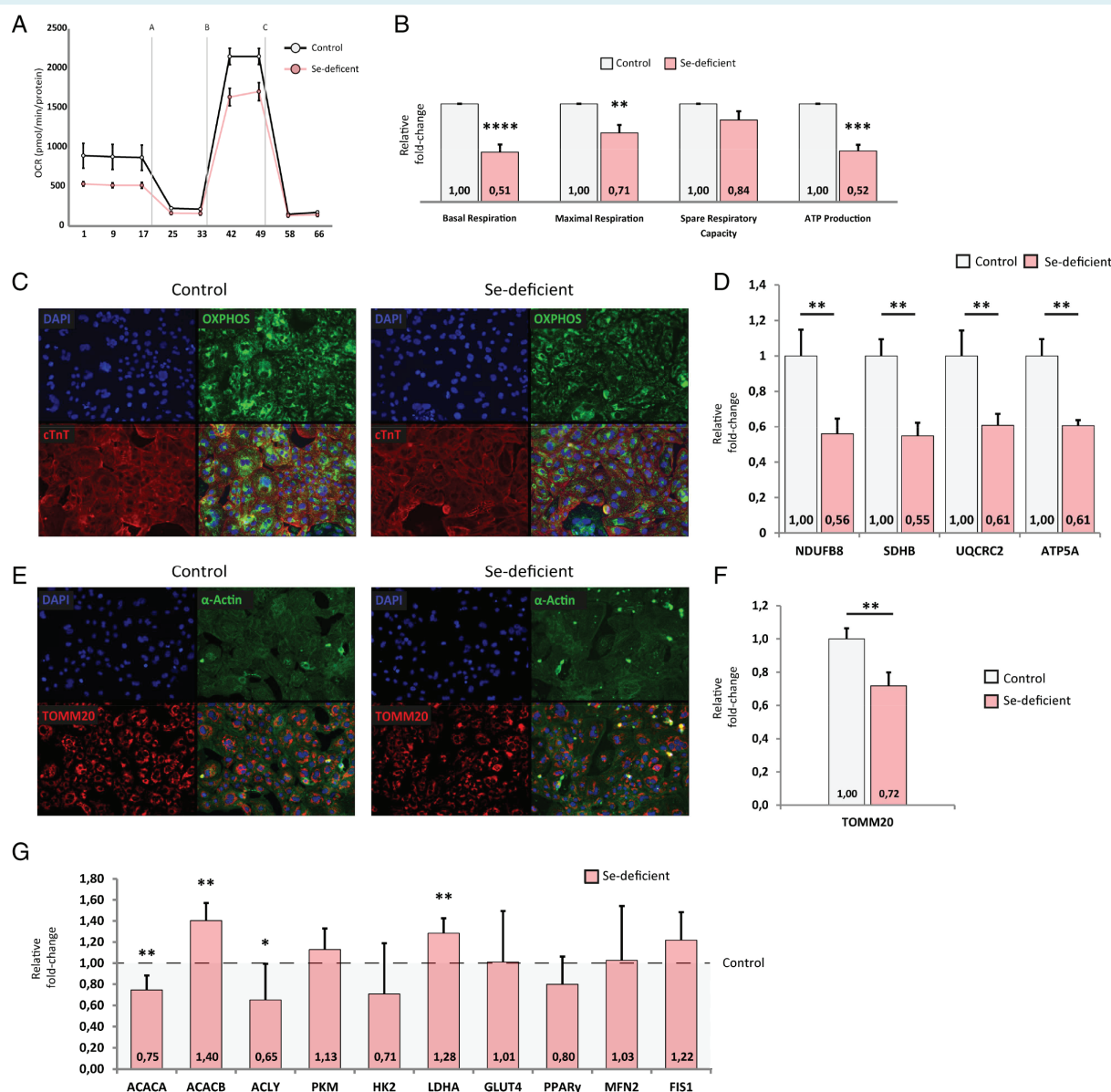


Figure 2 Effects of selenium (Se) depletion on mitochondrial respiration and metabolic function. (A) Representative traces for control cardiomyocytes and Se-deficient cardiomyocytes in a Mito Stress Test. (B) Effects of Se deficiency on basal respiration, maximal respiration, respiratory reserve and ATP-linked respiration ($n = 5$). Representative immunofluorescent staining for (C) DAPI, OXPHOS and cardiac troponin T (cTnT) and (E) DAPI, TOMM20 and α -actin in control and Se-deficient cardiomyocytes ($n = 3$). Western blot quantification of (D) NDUF8, SDHB, UQCRC2 and ATP5A (mitochondrial complexes 1 to 3 and 5, respectively) ($n = 6$) and (F) TOMM20 ($n = 6$). (G) Relative fold-change of genes involved in (anaerobic) glycolysis or fatty acid metabolism and mitochondrial fusion and fission ($n = 6$). OCR, oxygen consumption rate. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$, **** $P < 0.0001$.

chain function leading to less efficient production of ATP, increased production of ROS and intracellular oxidative damage. However, selenium may not only be a marker of greater disease severity and worse outcome but also a therapeutic target. We found that the relationship between selenium deficiency and signs, symptoms and prognosis were similar or even more pronounced than for iron deficiency. Clinical trials of intravenous iron supplements have shown improvements in symptoms and well-being, and trials

investigating effects on morbidity and mortality are underway. Unlike iron, selenium is readily absorbed orally. Trials of selenium supplementation are now required.

Selenium accumulates in biological tissue and is generally known as an antioxidant due to its presence in selenoproteins as selenocysteine. Furthermore, it is known to affect production of active thyroid hormone T_3 ²⁶ and anti-inflammatory and anti-apoptotic capacity.²⁷ With the occurrence of modest

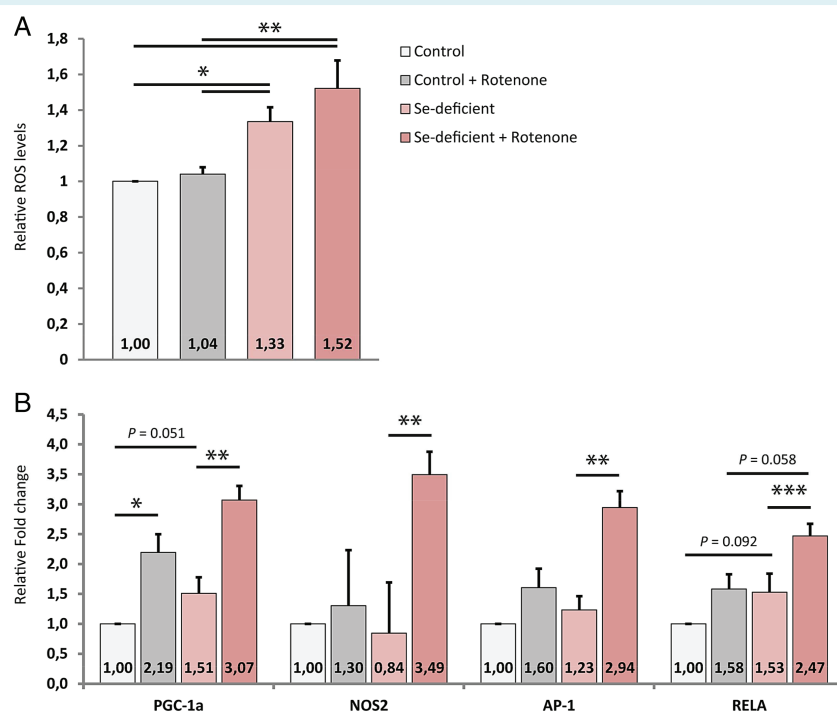


Figure 3 Effects of selenium (Se) depletion on reactive oxygen species (ROS) levels and transcriptional stress markers. (A) Relative ROS levels in control and Se-deficient cardiomyocytes, either with or without the addition of rotenone ($n = 5$). (B) Relative fold-change of genes involved in cardiomyocyte stress response upon oxidative stress ($n = 5$). * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

deficiency, selenoprotein concentrations and activities are preferentially lost based on their demand.²⁸ Serum selenium concentrations are reduced in many other age-related diseases, including cancer and heart failure.^{7,28} Appetite and therewith food intake may be reduced because of dietary advice on salt and calorie intake, social isolation, possible circulatory congestion and inflammation-induced malabsorption, although evidence on the latter is conflicting.^{12,29} Age-matched healthy controls showed higher selenium levels than patients with worsening heart failure (online supplementary Table S3), with only 5% of controls $<70 \mu\text{g/L}$. Selenium toxicity can occur with ingestion of excess selenium and symptoms include nausea, vomiting, nail brittleness and loss, hair loss, fatigue, irritability and foul breath odour as found in a study investigating a misformulated liquid dietary supplement that resulted in 201 cases of selenium poisoning [the median estimated amount of selenium ingested was 989 mg (range: 41–5875 mg)].³⁰ Nonetheless, selenium supplementation is considered to be safe with a recommended dietary allowance of 55 $\mu\text{g/day}$ for persons ≥ 14 years, with a tolerable upper intake limit of 400 $\mu\text{g/day}$.³⁰

A meta-analysis of 25 observational studies on selenium supplementation showed a moderate association between increasing serum selenium concentration and reduced risk of coronary heart disease.¹⁰ However, randomised trials of selenium supplementation did not show a profound reduction in cardiovascular events or mortality,¹⁰ but these were often performed in small cohorts and contain several confounding factors (e.g. age and baseline selenium levels) hampering interpretation and generalization. One major

confounding factor in these trials is the high mean serum selenium levels, especially in cohorts including subjects from countries with relatively high intake (USA, Canada).³¹ Beyond a specific selenium concentration, there may be no further advantage of selenium supplementation in reducing cardiovascular mortality.³² This is supported by our data on exercise capacity and quality of life, showing only for selenium concentrations $<100 \mu\text{g/L}$ a negative effect. Furthermore, selenium is often given in combination with other vitamins or minerals (e.g. zinc, vitamin C and E and β -carotene) making it difficult to identify selenium specific effects.¹⁰ A recently reported trial of older people (aged 70–80 years), many of whom were not known to have serious cardiac disease, randomly assigned participants to placebo or supplements of selenium 200 $\mu\text{g/day}$ and coenzyme Q10 for a median of 5.1 years.^{33,34} Daily selenium supplements increased serum selenium concentrations from 45–87 $\mu\text{g/L}$ at baseline to 185–245 $\mu\text{g/L}$ at 48 months, whereas it did not change in those assigned to placebo.³³ Those receiving supplements had lower plasma concentrations of NT-proBNP at 24 and 48 months and improved echocardiographic function.³⁴ Furthermore, participants with a serum selenium concentration $<65 \mu\text{g/L}$ had a higher cardiovascular mortality compared to those with concentrations $>85 \mu\text{g/L}$.³³ It is unclear whether there is a synergistic effect of selenium and coenzyme Q10 supplements. No randomized trial has evaluated the effect of selenium supplements alone in patients with heart failure.

The effects of selenium deprivation was assessed for the first time in human PSC-derived cardiomyocytes and showed to impair

mitochondrial function, biogenesis and oxidative stress. Mitochondria of selenium-deprived cardiomyocytes had a two-fold lower basal oxygen consumption rate and ATP-linked respiration. Furthermore, on gene expression level, we found decreased expression *ACACA* and *ACLY*, but increased expression of *ACACB* indicating decreased *de novo* lipogenesis and increased inhibition of fatty acid and glucose oxidation, respectively. *LDHA* was up-regulated as the result of low selenium, indicative of anaerobic glycolysis. This was in concordance with lower levels of oxidative phosphorylation, implicating a more glycolytic metabolism profile similar to failing cardiomyocytes.³⁵ We found no evidence for induced mitochondrial fission or fusion. One potential mechanism of benefit from selenium supplementation is the reduction of oxidative stress. ROS levels were increased in selenium-depleted cardiomyocytes under normal culture conditions. Upon inhibiting mitochondrial complex I, ROS levels increased even more with low selenium status, but not in cardiomyocytes with a sufficient selenium status, implying non-optimal working antioxidant machinery. In the acute phase following myocardial infarction and especially subsequent reperfusion, oxidative damage is predominant. One major source for myocardial ROS lies within the mitochondria and its production is linked to glutathione.^{33,36,37} Glutathione is a selenium-containing enzyme and an important antioxidant in human. Among the 25 selenoproteins in mammals, the family of glutathione peroxidases is a major group whose function is well understood.¹ Nevertheless, much is still unknown of the effect of the other selenoproteins and more research regarding the fundamental mechanisms of these proteins on cardiac homeostasis is necessary.

Limitations

In this retrospective study we only investigated the associations of selenium serum levels with clinical characteristics and outcomes. No data were currently available on other proteomic markers of selenium-dependent pathways, nor oxidative stress. As a result, other signalling components such as selenoprotein P or glutathione peroxidase were not investigated next to serum selenium. Furthermore, reference ranges for serum selenium vary between countries because of differences in dietary sources of selenium. For Caucasians of European ancestry, adult reference ranges are between 70 and 130 µg/L. Deficiency of selenium was therefore set at serum levels <70 µg/L. Finally, although numerous associations were identified between selenium and clinical measurements/outcomes, more research regarding the fundamental mechanisms of selenium on cardiac homeostasis and the potential benefits of selenium supplements in patients with heart failure and low serum selenium concentrations is necessary.

Conclusions

In conclusion, selenium is required to maintain health including normal mitochondrial and cardiac function. Many older people, with and without heart failure, have low serum concentrations of selenium, suggesting deficiency. In patients with heart failure, low serum concentrations are associated with more severe symptoms and signs and a worse prognosis. Selenium supplements safely

correct low serum concentrations and in a trial of older people, many of whom did not have serious cardiovascular disease, a combination of selenium and coenzyme Q10 supplements improved outcome. Selenium is readily available, inexpensive and has been thoroughly tested for toxicity, making it a potentially important 'nutraceutical'. Having said this, we would like to state that by using an observational study we are unable to attribute cause and effect of selenium deficiency. Although we show significant effects on mitochondrial function and ROS of *in vitro* cultured cardiomyocytes, these functional data are suggestive and do not constitute such proof. Therefore, therapeutic trials are needed to achieve conclusive evidence of a causative role of selenium deficiency in heart failure patients and to determine the benefits of selenium supplements in patients with heart failure and low serum selenium concentrations.

Supplementary Information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Methods S1. Supplementary methods.

Table S1. Primer sequences as used for RT-qPCR expression analyses.

Table S2. Multivariable logistic regression analysis between the prevalence of selenium deficiency (<70 µg/L) and baseline characteristics.

Table S3. Baseline characteristics of healthy controls against BIOSTAT-CHF patients.

Figure S1. Geographic variation in serum selenium concentrations.

Figure S2. Kaplan–Meier plots by quartile of selenium concentration for the primary endpoint of all-cause mortality or heart failure hospitalization and the secondary endpoint of all-cause mortality.

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